

## Product datasheet for **RC402738**

### Glypican 3 (GPC3) (NM\_004484) Human Mutant ORF Clone

#### Product data:

Product Type:	Mutant ORF Clones
Product Name:	Glypican 3 (GPC3) (NM_004484) Human Mutant ORF Clone
Mutation Description:	R387X
Affected Codon#:	387
Affected NT#:	1159
Nucleotide Mutation:	GPC3 Mutant (R387X), Myc-DDK-tagged ORF clone of Homo sapiens glypican 3 (GPC3), transcript variant 2 as transfection-ready DNA
Effect:	Simpson-Golbi-Behmel syndrome
Symbol:	GPC3
Synonyms:	DGSX; GTR2-2; MXR7; OCI-5; SDYS; SGB; SGBS; SGBS1
E. coli Selection:	Kanamycin (25 ug/mL)
Mammalian Cell Selection:	Neomycin
Vector:	pCMV6-Entry (PS100001)
Tag:	Myc-DDK
ACCN:	NM_004484
ORF Size:	1158 bp
Restriction Sites:	Sgfl-Mlul



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**ORF Nucleotide Sequence:**

>RC402738 representing NM\_004484  
 Red=Cloning site Blue=ORF Green=Tags(s)

TTTTGTAATACGACTCACTATAGGGCGGCCGGGAATTCGTCGACTGGATCCGGTACCGAGGAGATCTGCC  
 GCC**CGGATCGCC**

ATGGCCGGGACCGTGCACCGCGTGTGGTGGTGGCGATGCTGCTCAGCTTGGACTTCCCGGGACAGG  
 CGCAGCCCCCGCCGCGCCGCGGACGCCACCTGTACCAAGTCCGCTCCTTCTCCAGAGACTGCAGCC  
 CGGACTCAAGTGGGTGCCAGAACTCCCGTGCCAGGATCAGATTTGCAAGTATGTCTCCCTAAGGCCCA  
 ACATGCTGCTCAAGAAAGATGGAAGAAAAATACCAACTAACAGCACGATTGAACATGGAACAGCTGCTTC  
 AGTCTGCAAGTATGGAGCTCAAGTTCTTAATTATTGAGAAATGCTGCGGTTTTCAAGAGGCCCTTTGAAAT  
 TGTTGTTCCGCATGCCAAGAATAACCAATGCCATGTTCAAGAACAATAACCAAGCCTGACTCCACAA  
 GCTTTTGAGTTTGTGGGTGAATTTTTCACAGATGTGTCTCTACATCTGGGTTCTGACATCAATGTAG  
 ATGACATGGTCAATGAATGTTTGACAGCCTGTTCCAGTCATCTATACCCAGCTAATGAACCCAGGCCCT  
 GCCTGATTCAGCCTTGGACATCAATGAGTGCCTCCGAGGAGCAAGACGTGACCTGAAAGTATTTGGGAAT  
 TTCCCCAAGCTTATTATGACCCAGGTTTCCAAGTCACTGCAAGTCACTAGGATCTTCCCTCAGGCTCTGA  
 ATCTTGGAATTGAAGTATCAACACAATACTGATCACCTGAAGTTCAAGTAAAGGACTGTGGCCGAATGCTCAC  
 CAGAAATGGTACTGCTCTTACTGCCAGGGACTGATGATGGTTAAACCCTGTGGCGGTTACTGCAATGTG  
 GTCATGCAAGGCTGTATGGCAGGTGTGGTGGAGATTGACAAGTACTGGAGAGAATACATTCTGTCCCTTG  
 AAGAACTTGTGAATGGCATGTACAGAATCTATGACATGGAGAAGTACTGCTTGGTCTCTTTTCAACAAT  
 CCATGATTCTATCCAGTATGTCCAGAAGAATGCAGGAAAGCTGACCACCACTATTGGCAAGTTATGTGCC  
 CATTCTCAACAACGCCAATATAGATCTGCTTATTATCCTGAAGATCTCTTTATTGACAAGAAAGTATTA  
 AAGTTGCTCATGTAGAACATGAAGAAACCTTATCCAGC

AG**CGGACCG**ACGCGTACGCGCCGCTCGAGCAGAACTCATCTCAGAAGAGGATCTGGCAGCAAATGATATCC  
 TGGATTACAAGGATGACGACGA TAAGGTTTAA

**Protein Sequence:**

>RC402738 representing NM\_004484  
 Red=Cloning site Green=Tags(s)

MAGTVRTACLVVAMLLSLDFPGQAQPPPPPDATCHQVRSFFQRLQPGLKWPETPVPGSDLQVCLPKGP  
 TCCSRKMEEKYQLTARLNMEQLLQSASMEKFLIIQNAAVFQEAFEIVVRHAKNYTNAMFKNYPSLTPQ  
 AFEFVGEFFTDVSLYILGSDINVDMMNELFDSLFPVIYQMLMNPGLPDSALDINECLRGARRDLKVFGN  
 FPKLIMTQVSKSLQVTRIFLQALNLGIEVINTDHLKFSKDCGRMLTRMWYCSYCQGLMMVKPCGGYCNV  
 VMQGCMAVVEIDKYWREYILSLEELVNGMYRIYDMENVLLGLFSTIHDSIQYVQKNAGLTTTIGKLC  
 HSQQRQYRSAYYPEDLFIDKKVLKVAHVEHEETLSS

SGP**TRRRLEQKLI**SEEDLAANDILDYKDDDDKV

**Restriction Sites:**

Sgfl-Mlul



**Gene Summary:**

Cell surface heparan sulfate proteoglycans are composed of a membrane-associated protein core substituted with a variable number of heparan sulfate chains. Members of the glypican-related integral membrane proteoglycan family (GRIPS) contain a core protein anchored to the cytoplasmic membrane via a glycosyl phosphatidylinositol linkage. These proteins may play a role in the control of cell division and growth regulation. The protein encoded by this gene can bind to and inhibit the dipeptidyl peptidase activity of CD26, and it can induce apoptosis in certain cell types. Deletion mutations in this gene are associated with Simpson-Golabi-Behmel syndrome, also known as Simpson dysmorphia syndrome. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Sep 2009]